EXHIBIT 12



PHARMACY MANAGEMENT CONSULTING SERVICES

Monthly Progress Report

To The

California Prison Health Care

Receivership Corporation

January 2007

Contents

Introduction	
Executive Summary of Key Points in this Report	. 2
Objectives Completed	. 2
Objectives Delayed	. 2
Obstacles or Issues for Success	. 2
Progress Report Detail by Goal	. 3
Goal A	
Actions Taken	. 3
Objectives Completed	. 5
Issues or Obstacles to Success	
Goal B	. 5
Actions Taken	. 5
Objectives Completed	. 6
Issues or Obstacles to Success	
Goal C	. 7
Actions Taken	
Objectives Completed	
Issues or Obstacles to Success	
Goal D.	. 9
Actions Taken	
Objectives Completed	
Issues or Obstacles to Success	
Goal E	
Actions Taken	. 9
Objectives Completed	10
Issues or Obstacles to Success	
Goal F	10
Actions Taken	10
Objectives Completed	
Issues or Obstacles to Success	
Goal G	11
Actions Taken	
Objectives Completed	11
Issues or Obstacles to Success	
Summary of Changes to Timeline	12
Objectives Completed	
Objective Timelines Proposed for Change	
Objective Timeline Change Approvals	
Conclusion	
Appendix A— Updated Timeline	13
Appendix B— Financial Summary	
Appendix C— Project Organization Chart & Authority Letter	
Appendix D— Pharmacy Dashboard	
Appendix E— Pharmacy Inspection Grid / Monthly Inspection Form / Baseline	
Inspection Profile	13

PHARMACY MANAGEMENT CONSULTING SERVICES

Monthly Progress Report January 2007

Introduction

The California Prison Health Care Receivership Corporation (CPR) and Maxor National Pharmacy Services Corporation (Maxor) entered into an agreement to provide pharmacy management consulting services to achieve necessary improvements to the California Department of Corrections and Rehabilitation (CDCR). The implementation of the Agreement made effective January 1, 2007, commenced on schedule. Key members of the Maxor team arrived in Sacramento on January 1, 2007 and established an administrative office at 428 J Street Ste 610, Sacramento, CA 95814.

With the approval of the Receiver, the Maxor team was able to recruit and make immediately available to the CPR experienced and well qualified correctional pharmaceutical clinicians to wit: Glenn Johnson, MD, Project Manager, Matt Keith, RPh, BCPS, FASHP, Pharmacy Administrator, Dick Cason, RPh, MS, Senior Pharmacy Consultant and Melanie Roberts, RPh, PharmD, Clinical Pharmacy Consultant. Collectively, the management consulting team has over 70 years of direct oversight involvement with correctional and commercial pharmacy programs nationwide.

During this reporting period, the Receiver's office was instrumental in transitioning the Maxor team into the CPR program. Breakout meetings were arranged, security clearances achieved, administrative and operational guidance given and most importantly, the Receiver's direction and priorities were established. These priorities include working closely with the Court's experts in the *Coleman* (mental health) and *Perez* (dental) litigation. Additionally, the Receiver requested and Maxor provided an initial 90 day plan of specific implementation actions which was subsequently reviewed and approved by the Receiver on January 24th.

The collective efforts of the pharmacy improvement program will evolve around the court approved *Road Map to Excellence* adopted by the CPR with priority given to achieving patient safety, evidenced based practice and cost efficiency. Emphasis will be placed on efforts to ensure effective *Plata/Coleman/Perez* interfaces. The required improvements outlined in the *Road Map* are organized into seven primary goals. Each goal is supported by specific objectives and timelines for accomplishing those objectives.



This document provides a status report of the progress made during the month towards achieving each goal, summarizes any changes to the projected timelines, identifies potential obstacles or issues that may delay or impact progress and provides an updated timeline (Appendix A) and financial status (Appendix B) for the project.

Executive Summary of Key Points in this Report

The following summary highlights key accomplishments, identifies any delays experienced and notes obstacles or issues related to achieving the required goals and objectives explained in more detail within this month's Progress Report.

Objectives Completed

- A central pharmacy services administration, budget and enforcement authority was established. (Objective A.1.)
- Direct lines of authority to all pharmacy services personnel and linkage to central medical staff were established with the Receiver's approval. (Objective A.2)

Objectives Delayed

• All objectives are progressing according to schedule.

Obstacles or Issues for Success

Several issues have been identified that may impact the achievement of planned objectives within the expected timeframes or require modification of scheduled work products. Each of these items is being addressed by the Maxor project team with the assistance of the Receiver.

- There is currently no active process for central operational procedure review and approval.
- CDCR lacks a central pharmacy information management system which has contributed to delays in collecting purchasing, prescription and outcomes data for process improvement. Current pharmacy utilization data are unreliable and restricted to purchases. Until such a system is in place, data will remain unreliable.
- The system-wide inventory that was to be conducted during the first quarter to establish a baseline would be of no value in inventory tracking in light of the absence of a CDCR pharmacy management system capable of accurately and uniformly recording dispensing data. Accordingly, Maxor is proposing a change in the timeline of Objective C.2.1. We request to limit the requirement of a system-wide inventory to controlled substances. A system-wide full inventory should occur with the deployment of an interim pharmacy system, which may occur in the first quarter, but is more likely to be completed in the second quarter.



- Initial delays were encountered when attempting to compare CDCR drug purchases with Department of General Services (DGS) contract prices. Access to one specific contract was delayed for a period of several weeks due to legal review.
- Although obtaining and disseminating documentation of Maxor's authority to intercept, review and credit orders to the wholesaler took longer than anticipated, a letter of authority from the Receiver's Chief of Staff was provided January 23, 2007 and actions are now underway.
- An initial audit of CDCR purchases and contract pricing since the initial 2006 Maxor review was conducted. A detailed listing of overcharges in the amount of \$299,000 has been sent to the wholesaler. Since November 2005, eligible rebates in the amount of \$474,000 for Zyprexa were reconciled and the receipt of \$343,000 has been confirmed. The additional \$131,000 is still being researched to ensure credit was received.

Progress Report Detail by Goal

For each goal in the *Road Map*, a summary of actions taken and progress achieved during the last 30 days is listed, along with the identification of any obstacles or issues that may impede progress.

Goal A

Develop meaningful and effective centralized oversight, control and monitoring over the pharmacy services program.

Actions Taken

- Maxor's Sacramento office was established on January 1, 2007 with the opening of project headquarters at 428 J Street Ste 610, Sacramento, CA 95814 (Objective A.1)
- Maxor project team members arrived on site for orientation and initial briefing by Corporate. The team includes a Project Manager, Senior Pharmacy Administrator and two pharmacy consultants. (Objective A.1; also see Appendix C)
- Meetings were conducted with the Receiver and staff to receive initial guidance and project direction. (Objective A.1)
- In conjunction with the Receiver's staff attorney, plans were developed and timelines established for implementing centralized oversight, control and monitoring over the CDCR pharmacy services program. (Objective A.1)
- An initial 90 day plan of action was prepared by the Maxor team and submitted for review and approval to the Receiver's Office. Subsequently, the 90-day plan was approved by letter dated January 24, 2007, from John Hagar, Chief of Staff to the Receiver. Maxor was also directed to work closely with the Court's experts in the additional two health care legal cases *Coleman* (mental health) and *Perez* (dental).



- Job descriptions for Director, Assistant Director and Clinical Pharmacists were finalized. (Objective A.1)
- Recruitment of the Pharmacy Director, Assistant Director and Clinical Pharmacy specialists commenced. Maxor is in the process of assessing and making an offer to an Assistant Director candidate. (Objective A.1)
- An agreement was established with the Receiver on the recruitment of 8 clinical pharmacists and a supplemental fiscal note was submitted for approval. This note establishes the clinical positions as Maxor employees. In addition, the agreement creates a "drop-in" strike team to include a manager and 4 technicians to help remedy problems in pharmacies with significant and immediate service issues. (Objective A.1)
- Early interaction by the Maxor team has additionally identified the need for a professional pharmacy nurse liaison to assist with nursing issues associated with distribution and administration of medications to patients and aid the Maxor Project Manager's efforts to ensure effective *Plata/Coleman/Perez* interfaces. A supplemental budget request was submitted by the Maxor team to the Receiver's Chief of Staff, and has been forwarded to the Receiver with a recommendation for approval. (Objective A.1)
- Meetings with CPR and CDCR staff have commenced and have included interactions with Dr. Terry Hill, Dr. Odegaar-Turner, Dr. Kuykendall, Linda Buzzini, John Hummell, Dr. Justin Graham, the DGS and AmeriSource Bergen to discuss the *Road Map* and future relationships. (Objective A.2)
- A clear organizational chart of reporting relationships and chains of command and coordination was developed with input from and approval by the Receiver's Chief of Staff. A letter was sent by the CPR Chief of Staff to CDCR officials delineating the role of Maxor. (Objective A.2; see Appendix C)
- Lines of communications were established with the CPR and CDCR Health Services office. (Objective A.2)
- With the approval of the Receiver, system wide CDCR Pharmacy staff information briefings on the *Road Map* were scheduled. (Objective A.2)
- A Regional Provider meeting was conducted on January 30, 2007 to provide orientation to the *Road Map* and the organizational structure established by Maxor and the Office of the Receiver. (Objective A.2)
- Central Policies & Procedures were obtained for review during the first quarter. A collection of facility specific Policies & Procedures are also being assembled. (Objective A.3)
- Early policy and procedure revisions will target the Pharmacy & Therapeutics Committee empowerment and essential activities. Suggested changes will be presented to the P&T Committee during its initial February 2007 meeting. (Objective A.3)
- A dashboard of pharmacy utilization was created and is being populated as data becomes available. Information sources and data reliability continue to be assessed. The existing pharmacy system data has been determined to be unreliable, and non-reproducible. Current data sources are limited to medication purchases without individual medical record review. Until a central pharmacy



- information management system is instituted, data will remain unreliable. (Objective A.4 & A.5; see Appendix D)
- A project management file structure was created to track activities and initiatives as they develop (Objective A.5)
- An initiative timeline & tracking grid was developed to monitor implementation of the *Road Map*. (Objective A.4 & A.5: see Appendix A)
- A stoplight grid and facility inspection tool was developed to assess baseline facility level adherence to regulations, standards and security concerns. (Objective A.5; see Appendix E)
- A schedule for completing facility inspections has been developed and will be deployed in February. (Objective A.5)
- A disease management guideline (asthma) is under development and will be presented for review of content and form at the March P&T Committee meeting. (Objective A.5)

Objectives Completed

- Objective A.1. A central pharmacy services administration, budget and enforcement authority was established on January 23, 2007.
- Objective A.2. A pharmacy organizational chart was approved by the Receiver establishing direct lines of authority to all pharmacy services personnel and defining linkage to central medical staff.

Issues or Obstacles to Success

- There is currently no active process for central operational procedure review and approval. Recommendations for changes in Pharmacy Policy & Procedures will be prepared for discussion at the first system-wide P&T Committee scheduled for February 13, 2007.
- Current pharmacy utilization data are unreliable and restricted to purchases. There is no reproducible or reliable system for tracking dispensing or outcomes data. Until a central pharmacy information management system is instituted, data will remain unreliable.

Goal B

Implement and enforce clinical pharmacy management processes including formulary controls, Pharmacy and Therapeutics committee, disease management guidelines, and the establishment of a program of regular prison institution operational audits.

Actions Taken

• A reconstituted P&T Committee is being formed—anticipated members are 3 physicians (recommended by Regional MDs), Dr. Odegaard-Turner (Nursing), Jacki Clark (Nursing), Dr. Kuykendall (Dental Services), Dr. Hill, Dr. Winslow, Dr. Jeff Metzner (Colemen psychiatric expert), Dr. Andrew Swanson (Psychiatry) and the Maxor team (Johnson, Keith, Roberts and Cason). (Objective B.1)



- The initial P&T meeting is scheduled for Tuesday, February 13, 2007. The schedule for future meetings will be decided on at the initial meeting. (Objective B.1)
- A P&T Committee Charter is to be established at the first meeting (Objective B.1)
- Health Services Policy & Procedure changes will be recommended to the P&T Committee to establish their organizational structure and empowerment. (Objective B.1)
- A routine agenda format for the P&T Committee has been developed and will be presented. (Objective B.1)
- A new Correctional Formulary (based on the California Common Drug Formulary) is under development and will be presented in February for P&T review and approval. (Objective B.l)
- Formulary adherence and compliance will be reviewed upon approval of the Correctional Formulary. However, it should be noted that until the pharmacy management system is in operation, data gathering will continue to be unreliable. (Objective B.2)
- Therapeutic category utilization reports are being created and trends analyzed for discussion at first P&T meeting. (Objective B.2)
- The first disease management guideline (asthma) will be presented for review of content and style in March. This guideline will serve as the prototype for developing additional guidelines. (Objective B.3)
- Therapeutic category utilization data are being reviewed to determine a schedule for guideline development. Guidelines will focus on chronic illnesses to include asthma, hypertension, diabetes, hyperlipidemia and seizure disorder. Additional guidelines will be created as scheduled by the P&T Committee. (Objective B.3)
- Facility audit procedures have been established. Initial audits will begin statewide in February 2007. (Objective B.4; see Appendix E)

Objectives Completed

• All objectives are in progress.

Issues or Obstacles to Success

CDCR lacks a central pharmacy information management system which has
contributed to delays in collecting prescription and outcomes data for process
improvement. Maxor has an interim system in place to collect purchasing and
dispensing data. Processing this data is cumbersome and data reliability is a
concern as research indicates that dispensing data are not always entered into the
existing systems. An integrated pharmacy information management system is
vital for all reporting functions in order to provide true targeted process
improvement.



Goal C

Establish a comprehensive program to review, audit and monitor pharmaceutical contracting and procurement processes to ensure cost efficiency in pharmaceutical purchases.

Actions Taken

- CDCR purchases were downloaded from Amerisource Bergen (wholesaler) and an audit of contract pricing since the last review was conducted. (Objective C.1)
- A detail listing of overcharges in the amount of \$299,000 has been sent to the wholesaler. A meeting was held to discuss the credit/re-bill process. (Objective C.1)
- Since November 2005, eligible rebates in the amount of \$474,000 for Zyprexa were reconciled and the receipt of \$343,000 has been confirmed. The additional \$131,000 is still being researched to ensure credit was received. A retrospective review from the beginning of the rebate contract through November 2005 will be conducted. (Objective C.1)
- Until a comprehensive data system is in place to intercept, review and edit orders submitted daily to ensure contract compliance, a review schedule has been put in place that continually monitors CDCR purchases to assure that the correct price is charged, eligible rebates are obtained, and contract terms are met. (Objective C.1)
- In light of the absence of a CDCR pharmacy system capable of accurately and uniformly recording dispensing data, we will request a modification to the scope and timing of Objective C2.1. Conducting a system-wide inventory in order to establish a baseline at this time would not be of value from an inventory tracking standpoint.
 - The scope change requested is to limit the requirement of a system-wide inventory and establishment of a baseline to Controlled Substances until an interim pharmacy system is implemented. Controlled Substances have manual dispensing logs that can be used to inventory and immediately begin tracking during the first quarter in accordance with the original time requirement. We will request that a system-wide full inventory occur in conjunction with the deployment of an interim pharmacy management system, which may occur in the first quarter, but is more likely to be completed in the second quarter. This would also allow an audited inventory to be initially loaded into the system from the very beginning. (Objective C.2)
- A procedure is in place to compare all purchases versus dispenses to identify potential diversions or misuse. A written inventory control procedure will be drafted in the first quarter once facility audits are completed. (Objective C.2)
- A meeting with the Department of General Services and the wholesaler was held in order to establish an arrangement for the coordination and improvement of pharmaceutical procurement and contracting activities. An agreed upon method for standardizing order entry through the Echo System was suggested. Once the orders are consolidated in Echo, Maxor's supply system will "intercept" the order,



review it for contract compliance in conjunction with availability at the wholesaler's local distribution centers, edit the order and resubmit it electronically. This will ensure that the best value contracted item is purchased and enable dialogue with the wholesaler to continually stock the appropriate contracted items necessary to meet the demands of CDCR. This process is not anticipated to create a delay in treatment or delivery of the requested drugs. (Objective C.3)

- DGS was provided with access to Maxor's enterprise reporting system and given instruction on how to use the tools to better monitor purchasing for contract compliance and future negotiations. (Objective C.3)
- A meeting with the CPR's Chief Medical Information Officer and Chief Information Officer was held to discuss interim solutions to immediately begin capturing uniform dispensing data and improve patient safety. An agreement will be made after a thorough review of available interim solutions and a CDCR pilot system (Pelican Bay) currently in place. (Objective C.3)
- A review of 340B pricing feasibility has been initiated. This study will assess the feasibility of achieving cost savings through the utilization of 340B pricing to mitigate the rising prescription drug expenditures by the CDCR. This study will be conducted to quantify the potential cost savings for California taxpayers resulting from access to 340B pricing by the CDCR. The study will also address the potential barriers associated with implementing this strategy and the initial steps necessary for establishing a 340B Drug Discount Program. A request to the Receiver's Chief of Staff for permission to engage the Heinz Family Foundation to assist in this review has been made. (Objective C.5)

Objectives Completed

All objectives are in progress.

Issues or Obstacles to Success

- Initial delays were encountered when attempting to compare CDCR drug purchases with DGS contract prices. Access to one specific contract was delayed for a period of several weeks due to legal review by DGS. On January 26, 2007, Maxor received a letter from the DGS Office of Legal Services expressing their intent to provide the requested contracts. On January 29, 2007, Maxor received the delayed contract in question.
- Although obtaining and disseminating documentation of Maxor's authority to intercept, review and credit orders to the wholesaler took longer than anticipated, a letter of authority was provided January 23, 2007 and actions are now underway.



Goal D

Develop a meaningful pharmacy human resource program that effectively manages staffing, compensation, job descriptions, competency, performance assessment, discipline, training, and use of the workforce including temporary employees and non pharmacist staff.

Actions Taken

- Maxor has requested all staffing agency service contracts for review. Billing summaries will also be reviewed and audited. (Objective D.1)
- Maxor is obtaining staffing levels and position descriptions (CDCR, Registry, Vacant) to populate an employee tracking system. The system will allow the Maxor Team to identify vacancies to be filled as well as provide a tracking mechanism for employee training, education and disciplinary actions. (Objective D.2)
- Web based software was reviewed to identify a product that could provide information (education & training modules created by the Maxor team) to CDCR pharmacy staff and allow for competency assessment tests. Test tracking will assure all staff complete required training. The products assure deployment of important procedural changes, educational information and other key information. (Objective D.2)

Objectives Completed

• All objectives are in progress

Issues or Obstacles to Success

No significant issues or obstacles encountered to date.

Goal E

Redesign and standardize overall institution level pharmacy drug distribution operations for inpatient and outpatient needs. Design, construct and operate a centralized pharmacy facility.

Actions Taken

- An assessment of potential sites for establishing a centralized pharmacy facility
 has commenced and including at a minimum: Fresno, Stockton and Sacramento.
 Criteria include access to lines of transportation (air and ground), location,
 proximity to pharmaceutical distribution centers, ability to recruit and maintain
 qualified pharmacy staff and costs. (Objective E.2)
- Contact with potential sources of prepackaged product is occurring. The outsourced product will be considered for pre-centralization to assist facilities in meeting their service and product control needs. (Objective E.2)



• A Maxor team member spent two days at San Quentin reviewing pharmacy and nursing practices and assessing immediate service needs. Operating procedures for a "drop-in" strike team are being formulated. (Objective E.1)

Objectives Completed

• All objectives are in progress

Issues or Obstacles to Success

• No significant issues or obstacles encountered to date.

Goal F

Based on a thorough understanding of redesigned work processes, design and implement a uniform pharmacy information management system needed to successfully operate and maintain the CDCR pharmacy operation in a safe, effective and cost efficient way.

Actions Taken

- A repository of prescription data from the existing PDTS system has been designed for consistent data accumulation and reporting. (Objective F.1)
- Rudimentary utilization reports have been designed and will be distributed to the Receiver and the P&T Committee on a monthly basis and electronically to facilities when connectivity is established. The reports will become more sophisticated as data collection becomes more reliable. (Objective F.1)
- The CPR's CIO agreed to address all of the connectivity issues between the institutions. In the interim, we will work with him to establish connectivity using commercial digital subscriber line (DSL) and virtual private network (VPN) solutions. (Objective F.2)
- A list of initial equipment necessary to modernize the institutions and implement an interim solution has been provided to the CIO for procurement/authorization. (Objective F.2)
- The use of *Guardian Rx*, a pharmacy information management system that Maxor uses extensively in several operations throughout the nation, is currently being evaluated as a possible interim solution. IT requirements have been identified and submitted to the Receiver's IT representatives. (Objective F.3)

Objectives Completed

All objectives are in progress.

Issues or Obstacles to Success

No significant issues or obstacles encountered to date.



Goal G

Develop a process to assure CDCR pharmacy meets accreditation standards of the designated health care review body (NCCHC or ACA) and assist in obtaining accredited status.

Actions Taken

• No action taken in the first 90-days, pending completion of related objectives.

Objectives Completed

• No objectives completed.

Issues or Obstacles to Success

• No issues or obstacles to date.

Summary of Changes to Timeline

In the sections below, a listing of completed objectives, objective timelines proposed for change (subject to review and approval of CPR) and a listing of timeline changes that have been approved by the CPR are provided.

Objectives Completed

- Objective A.1. A central pharmacy services administration, budget and enforcement authority was established on January 1, 2007.
- Objective A.2. Received Receiver's approval to establish direct lines of authority to all pharmacy services personnel and define linkage to central medical staff.

Objective Timelines Proposed for Change

- As discussed under Goal C above, Maxor requests approval of a modification to the scope and timing of Objective C.2.1:
 - o The scope change requested is to limit the requirement of a system-wide inventory and establishment of a baseline to Controlled Substances until an interim pharmacy system is implemented.
 - O Modification of the timeline to reflect that a system-wide full inventory will occur in conjunction with the deployment of an interim pharmacy system, which may occur in the first quarter, but is more likely to be completed in the second quarter.

Objective Timeline Change Approvals

• Objective C.2.1 - timeline change approval pending.

Conclusion

Maxor remains committed to the accomplishment of the *Road Map* goals and objectives and has prepared this Progress Report as part of its ongoing initiative to maintain direct, open and constant communication with CPR throughout the pharmacy improvement project.

Maxor would like to thank the Receiver, his staff, and CDCR for their cooperation and support.



Appendix A— Updated Timeline

Appendix B— Financial Summary

Appendix C— Project Organization Chart & Authority
Letter

Appendix D— Pharmacy Dashboard

Appendix E— Pharmacy Inspection Grid / Monthly
Inspection Form / Baseline Inspection
Profile

APPENDIX A

Maxor Timeline and Tracking Grid for Accomplishing Roadmap Objectives

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- (1) The timeframes are contingent upon prerequisite approvals, funding and regulatory issues being addressed in a timely manner.

 (2) Some activities may begin earlier than shown and other activities may slide forward dependent upon the completion of related activities.
- (3) Ongoing activities may include addressing any lingering implementation issues, as well as addressing the transition of activity to the CDCR.
- (4) A proposed progress report schedule is included for documenting the accomplishments and identifying the need for schedule changes.

APPENDIX C

CPR Office of the California Prison Receivership Robert Sillen, Receiver

Receiver's San Francisco Office Federal District Courthouse Law Library 18th Floor 450 Golden Gate Avenue San Francisco, CA 94102

January 14, 2007

To: Wardens

Health Care Managers/Chief Medical Officers Regional Administrators Regional Directors of Nursing Pharmacists in Charge

Effective January 1, 2007 the Maxor National Pharmacy Services Corporation ("Maxor") commenced a contract with the California Prison Health Care Receivership Corporation ("CPR") to provide pharmacy management consulting services to CPR. Concerning these services, Maxor functions as a consultant empowered by the Receiver to perform services, including direct management services, as summarized in the "Maxor Timeline and Tracking Grid for Accomplishing Roadmap Objectives." In this regard, the requirements for cooperation with the Receiver set forth in Judge Henderson's Order of February 14, 2006 apply fully to Maxor.

Maxor is in the process of establishing direct lines of authority over all California Department of Corrections and Rehabilitation pharmacy services personnel, and is also in the process of establishing direct relations with California's control agencies, including but not limited to the Department of Finance and the Department of General Services. Maxor has established an office at 428 J Street, Suite 610, Sacramento California, 95814 and Dr. Glenn Johnson, Maxor's Project Manager can be reached by telephone at (916) 441-1089.

Yours truly,

John Hagar Chief of Staff

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ASP - Avenal State Prison
CAL - Calipatria State Prison
CCC - Ca Corr Center CIM - Ca Institute for Men
CIW - Corr Institute for Women CMC - Ca Men's Colony CCWF - Central Ca Women's Facility VSPW - Valley State Prison for Women WSP - Wasco State Prison SVSP - Salina Valley State Prison SATF - California Substance Abuse TF

C - Sierra Conservation Center PVSP - Pleasant Valley State Prison RJD - RJ Donovan Corr Facility CEN - Centinela State Prison CCI - Ca Corr Institute SQ - San Quentin SAC - California State Prison, Sacramento MCSP - Mule Creek State Prison
NKSP - North Kern State Prison
PBSP - Pelican Bay State Prison ISP - Ironwood State Prison
KVSP - Kern Valley State Prison DVI - Deuel Vocational Institute CVSP - Chuckawalla Valley State Prison CMF - Ca Medical Facility CMC - Ca Men's Colony CAL - Calipatria State Prison HDSP - High Desert State Prison CRC - Ca Rehabilitation Center COR - Ca State Prisons, Corcoran CEN - Centinela State Prison CCWF - Central Ca Women's Facility CCI - Ca Corr Institute CCC - Ca Corr Center ASP - Avenal State Prison WORKLOAD AC - Ca State Prison LA TF - CorrTraining Facility L - Ca State Prison, Solano 1 - Ca Institute for Men Staffing N - Corr Institute for Women Measure CDCR Average Rx #/Pharmacy **Measure Definitions** RPH# RX # Mo Avg Mo Avg CY 2005 Mo Avg Mo Avg Actual CY 2006 Mo Avg Mo Avg 45 GE Jan-07 Feb-07 Mar-07 Apr-07 May-07 Jun-07 Jul-07 Aug-07 Sep-07 Oct-07 Nov-07 Dec-07 Target or FY06 vs FY07 Stoplight Status (R/Y/G) Detail Data

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PBSP - Pelican Bay State Prison DVI - Deuel Vocational Institute CMF - Ca Medical Facility
COR - Ca State Prisons, Corcoran MCSP - Mule Creek State Prison ISP - Ironwood State Prison

KVSP - Kern Valley State Prison CRC - Ca Rehabilitation Center CVSP - Chuckawalla Valley State Prison WORKLOAD VSP - Pleasant Valley State Prison TF - CorrTraining Facility AC - Ca State Prison LA)L - Folsom C - Ca Corr Center JSP - High Desert State Prison Staffing . - Calipatria State Prison Measure CDCR Average RPh/Pharmac Measure Definitions Tech # ₽ * Mo Avg Mo Avg CY 2005 Mo Avg Mo Avg Actual CY 2006 Mo Avg Mo Avg 2007 TD Jan-07 Feb-07 Mar-07 Apr-07 May-07 Jun-07 Jul-07 ellow - Short of target Aug-07 Sep-07 Oct-07 Nov-07 Dec-07 Target or FY06 vs FY07 Stoplight Status (R/Y/G) Detail

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NKSP - North Kern State Prison COR - Ca State Prisons, Corcoran
CRC - Ca Rehabilitation Center
CTF - CorrTraining Facility HDSP - High Desert State Prison DVI - Deuel Vocational Institute /SPW - Valley State Prison for Women PBSP - Pelican Bay State Prison FOL - Folsom CVSP - Chuckawalla Valley State Prison CMF - Ca Medical Facility CMC - Ca Men's Colony CIW - Corr Institute for Women CIM - Ca Institute for Men CEN - Centinela State Prison SVSP - Salina Valley State Prison SQ - San Quentin SCC - Sierra Conservation Center SP - Kern Valley State Prison CCWF - Central Ca Women's Facility CCI - Ca Corr Institute CCC - Ca Corr Center CAL - Calipatria State Prison VSPW - Valley State Prison for Women SP - Wasco State Prison SOL - Ca State Prison, Solano SATF - California Substance Abuse TF ASP - Avenal State Prison WORKLOAD C - Ca State Prison LA Measure CDCR Average Rx/Tech CDCR Average Rx/RPh Measure Definitions Number of Rx/Tech Mo Avg Rx # Mo Avg CY 2005 Mo Avg Mo Avg CY 2006 Mo Avg Mo Avg 2007 710 Jan-07 Feb-07 Mar-07 Apr-07 May-07 Jun-07 Jul-07 ellow - Short of target Aug-07 Sep-07 Oct-07 Nov-07 Dec-07 FY06 vs FY07 Stoplight Status (R/Y/G) Detail Data

Jerry Hodge, RPh, Chairman, Maxor Operational Support Rick Pollard Jim Riley, Sr. VP Correctional Healthcare John Ward, CEO Maxor California Department of General Services **APPENDIX C** (Eight) **Clinical Pharmacists** Regional Pharmacist Facility Pharmacies Clinical Pharmacy M. Roberts, PharmD Glenn Johnson, MD Man Keith, RPh, BCPS, FASHP Project Manager Pharmasy Administrator Asst. Director Regional Pharmacist Pharmacy (Clinical) Director Pharmacy Services California Prison Receivership **Pharmacists** CDCR Central Office Regional Pharmacist Sr. Pharmacy Consultant Dick Cason, RPh, MS **Pharmacy Operations** Central Fill Facility Division CDCR- Health Services Improvement Project CDCR Pharmacy Legend CDCR CPR **Maxor Project Team** Pharmacy Nurse Liaison* *Subject to CPR Approval COLUM K. Cloutier, RN

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